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22578-005US1

Application No.

10/535,345

Applicant

Semple, et al.

Filing Date

February 15, 2006

Group Art Unit
February 15, 2006

	U.S. Patent Documents						
Examiner Initial	Desig. ID	Document Number	Publication Date	Patentee	Class	Subclass	Filing Date If Appropriate
	AA						

	Foreign Patent Documents or Published Foreign Patent Applications							
Examiner	Desig.	,	Publication	Country or			Trans	lation
Initial	ID	Document Number	Date	Patent Office	Class	Subclass	Yes	No
	AB	WO06/069242A2	06/29/06	WIPO				
	AC	WO05044816A1	05/19/05	WIPO				
	AD	WO04103370A1	12/02/04	WIPO				
	AE	WO03078409A1	09/25/03	WIPO				
	AF	WO03062200A2	07/31/03	WIPO				
	AG	WO03022814A1	03/20/03	WIPO				
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	AN	EP1305286B1	12/08/04	Europe				
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	AP	Alterman, M. et al., "Fast microwave-assisted preparation of aryl and vinyl nitriles and the corresponding tetrazoles from organo-halides", J. Org. Chem. 65:7984-89 (2000)(supporting information attached)		
· ·	AQ	Cahn, R.S. et al., "Specification of molecular chirality", Angew. Chem. Internat. Edit. 5(4):385-415 (1966)		
	AR	Carballo-Jane et al., "Comparison of rat and dog models of vasodilation and lipolysis for the calculation of a therapeutic index for GPR109A agonists," <i>Journal of Pharmacological and Toxicological Methods</i> , Article in Press, doi:10.1016/j.vascn.2007.05.007 (2007).		
	AS	Carballo-Jane et al., "Comparison of rat and dog models of vasodilation and lipolysis for the calculation of a therapeutic index for GPR109A agonists," <i>Journal of Pharmacological and Toxicological Methods</i> , 56(3). pp. 308-316, (2007).		

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Information Discle by App		Applicant Semple, et al.		
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	вн	Katritzky, A. et al., "Alpha-lithiation of N-alkyl groups in pyrazoles", Tetrahedron Letters 39:2023-9 (1983)
	BI	Latli, B. et al., "Novel and potent 6-chloro-3-pyridinyl ligands for the &\text{\alpha}2 neuronal nicotinic acetylcholine receptor", J. Med. Chem. 42:2227-34 (1999)
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	BK	Maciejewski-Lenoir et al., "Langerhans cells release prostaglandin D <sub>2</sub> in response to nicotinic acid," Journal of Investigative Dermatology, 126:2637-2646 (2006).
	CA	Mahley, R. et al., "Drug therapy for hypercholesterolemia and dyslipidemia", Goodman & Gilman 36:971-1002
	СВ	Mariano, P. et al., "Mechanistic aspets of gas-phase photodecarbonylation reactions of bicycle[3.1.0]hexanones", J. Org. Chem. 45:1753-62 (1980)
	СС	Miller, R.D. et al., "Deoxygenation of sulfoxides promoted by electrophilic silicon reagents: preparation of aryl-substituted sulfonium salts", J. Org. Chem. 53:5571-3 (1988)
	CD	Movassaghi, M. et al., "A direct method for the conversion of terminal epoxides into $\gamma$ -butanolides", J. Am. Chem. Soc. 124(11):2456-7 (2002)
	CE	Newman-Evans, R. et al., "The influence of intramolecular dynamics on branching ratios in thermal rearrangements", J. Org. Chem. 55:695-711 (1990)
	CF	Nishimura, J. et al., "A novel synthesis of methylcyclopropanes", Tetrahedron Letters 25:2647-59 (1969)
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	СН	Prelog, V. et al., "Basic principles of the CIP-system and proposals for a revision", Angew. Chem. Int. Ed. Engl. 21:567-83 (1982)
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	CK	Semple et al., "Recent progress in the discovery of niacin receptor agonists," Current Opinion in Drug Discovery & Development, 10:452-459, (2007).		
	CL	Semple et al., "1-Alkyl-benzotriazole-5-carboxylic acids are highly selective agonists of the human orphan G-protein-coupled receptor GPR109b," <i>Journal of Medicinal Chemistry</i> 49:1227-1230, (2006).		
	СМ	Semple, "Niacin receptor agonists," <u>Presentation</u> , American Chemical Society 233 <sup>rd</sup> National Meeting & Exposition, March 25, 2007 – March 29, 2007, Chicago, Illinois		
	CN	Semple, "Discovery of selective agonists for GPR109a and GPR109b, the high and low affinity receptors for niacin," <u>Presentation</u> , <i>GPCRs in Medicinal Chemistry</i> , jointly organized by the Society of Chemical Industry, Royal Society of Chemistry and the Societa Chimica Italiana, September 18, 2006 – September 20, 2006, Verona, Italy		
	СО	Skinner et al, "Fluorinated pyrazole acids are agonists of the high affinity niacin receptor GPR109a,"  Poster, 30th National Medicinal Chemistry Symposium, June 25, 2006 – June 29, 2006, Seattle, WA		
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	CQ	Taber, D. et al., "Synthesis of the eight enantiomerically pure diastereomers of the 12-F2-Isoprostanes", J. Am. Chem. Soc. 124:13121-6 (2002)(supporting information attached)		
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	DA	Yagi, H. et al., "Removal of benzyl-type protecting groups frompeptides by catalytic transfer hydrogenation with formic acid", J. Org. Chem. 44(19):3442-4 (1979)		
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